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**ECONOMICAL EVALUATION OF THE CARBONATION-CALCINATION  
SYSTEM FOR CAPTURING CO<sub>2</sub>**

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**Abstract**

The cost of the CO<sub>2</sub> capture using carbonation-calcination cycles as a technological option of the CO<sub>2</sub> capture in post combustion is the sum of several terms: sorbent costs, capital costs related to the capture plant, the new operation and maintenance processes, the cost of the additional fuels necessary to compensate the penalty in the efficiency due to the capture, and the cost of the CO<sub>2</sub> compression.

The required conditions for the sorbents are that they have to be either very good or very cheap. The cost of the sorbent flow can't be higher than 1-3 €/per separated CO<sub>2</sub> ton. In the case of utilisation of CaCO<sub>3</sub> as a sorbent, it has been calculated that the cost per separated CO<sub>2</sub> ton is 2,5 €. The problem of this sorbent is that its deactivation is quick, but this is compensated by its high abundance in the nature.

Apart from this, it has to be into account other terms in order to obtain the global cost. It has supposed a generation plant of 360 MW with a fluidized bed reactor. It has been estimated that it can be reached a realistic capture up to 78% in the carbonator using

CaO as a sorbent, as well as a 100% of the CO<sub>2</sub> generated in the calcinator, resulting capture rates of 85% approximately.

The assumptions for the operation and maintenance are that the carbonators require a molar relation Ca/C of 4 for an effective CO<sub>2</sub> capture, a 30% of the fuel is burned in the calcinators and the rest in the combustor, the capacity of the CaO to capture CO<sub>2</sub> decreases with the number of cycles, the recuperation rate of CaO is 92,5%, the global efficiency of the plant is 40,7%, the fuel is petroleum coke or anthracite.

Furthermore, taking also into account the capital costs and the compression of the CO<sub>2</sub>, it has been calculated that the global cost is 15,5 €for a life of 30 years of the plant.

Therefore, a priori it seems to be a reasonable cost compared with the technological options of CO<sub>2</sub> capture that are in operation nowadays, specially those which use amines.

## **Introduction**

The CO<sub>2</sub> capture represents nowadays one of the most important options that are planned in order to decrease the emissions of this gas related to the combustion of the fossil fuels and to fight in this way the climate change

One of the existent solutions for capturing CO<sub>2</sub> is through the carbonation-calcination cycles, which is one of the technological options of CO<sub>2</sub> capture in post combustion, that consists in the CO<sub>2</sub> separation at a high temperature.

For this technology would be successful, it must be demonstrated its viability either technical or economical at an industrial scale. This last aspect is essential, and is for that reason that it has been realized a detailed economical analysis of this process.

## Economical Analysis

Making the economical analysis, it has been taken into account all the factors that determine the final price of CO<sub>2</sub> capture by means of this technology.

The cost of the CO<sub>2</sub> capture is the addition of several terms, where are included the sorbent costs, the elevated capital costs related to the capture plant, the new operation and maintenance costs, the cost of the additional fuels necessary to compensate the penalty in the efficiency due to the capture, and the CO<sub>2</sub> compression cost.

Firstly, an analysis about the sorbent costs has been realized. The CO<sub>2</sub> capture systems are relevant due to the elevated sorbent quantities that are used, at least high enough to equalize the CO<sub>2</sub> molar flow that is processed in the plant.

In our case, we have the limestone as sorbent (CaCO<sub>3</sub>), which has been compared with the systems based on amines (MEA), because they are the most used up to now in the industry. The requested condition to the sorbents is either very goods or very cheaps.

A low comparative cost in the composition of the sorbent per CO<sub>2</sub> unit separated, will be essential for any sorption-desorption system for capturing CO<sub>2</sub>. We can define the sorbent cost per CO<sub>2</sub> kg separated (COS) according to the following equation:

$$COS = \left( \frac{F_0}{F_R} \right) \left( \frac{F_R}{F_{CO_2}} \right) \frac{bM_s}{M_{CO_2}} C_s = \left( \frac{F_0}{F_R} \right) \frac{C_s^*}{M_{CO_2}}, \text{ where:}$$

- $COS$  : cost of sorbent makeup per tonne of CO<sub>2</sub> removed (€/kg of CO<sub>2</sub>)
- $F_0$  : makeup flow of sorbent (mol/s)
- $F_R$  : flow of sorbent required to react with the flow of CO<sub>2</sub> (mol/s)
- $F_{CO_2}$  : flow of CO<sub>2</sub> removed by the capture system (mol/s)
- $b$  : moles of sorbent required to stoichiometrically react with 1 mol of CO<sub>2</sub> (or O<sub>2</sub>)
- $M_s$  : molecular weight of the sorbent (kg/mol)
- $M_{CO_2}$  : 0,044 kg/mol
- $C_s$  : unit cost of sorbent (€/kg)

The term  $C_s^* = M_s C_s b \left( F_R / F_{CO_2} \right)$  is a suitable variable to compare the cost of the different sorbents, because it represents the cost of the quantity of sorbent required to react (or absorb) with 1 mol of CO<sub>2</sub> (or O<sub>2</sub>).

For a typical system selected as a reference and based on MEA,  $b = 2$ ,  $M_{MEA} = 0,061$ ,  $C_s = 1,35$  €/kg MEA, and the flow of the fresh sorbent is 1,5 kg of MEA per CO<sub>2</sub> tonne. The other terms correspond to values of  $F_0/F_R = 0,000152$  and  $F_R/F_{CO_2} = 3,57$ . Therefore, the term  $C_s^* = 0,589$  €/mol of CO<sub>2</sub> and a sorbent composition cost for systems at high scale based on MEA of approximately  $COS_{MEA} = 0,00203$  €/kg CO<sub>2</sub>, or 2,03 €/CO<sub>2</sub> captured tonne.

The logical basis that resides in the economical analysis is that the performance of the sorbent in any new system of CO<sub>2</sub> capture would be of an enough quality that assure that the fresh sorbent costs would be comparable with those that are being reached nowadays for a commercial technology already tested, for example, absorption with MEA. This is assumed to be a necessary condition (but not enough), because it is considered improbable that other benefits which come from a new and not tested technology could compensate the costs of the CO<sub>2</sub> capture process. However, one condition could be benefit for the introduction of such improvements, and this is that the emission of the CO<sub>2</sub> to the atmosphere was more expensive than its own capture.

For the regenerative sorbents, the energetic requirements represent the main contribution in the global capture costs.

The natural limestones have been studied as regenerable sorbents for the CO<sub>2</sub> separation. Nevertheless, the sorbent cannot maintain a high capacity capture further than 20 cycles, and are required high quantities of limestone for the CO<sub>2</sub> control. This only can be accepted because the low price of the limestone and because the calcinated products could acquire a value for the cement industry.

If we analyze the necessary sorbent for the CO<sub>2</sub> capture by means of the carbonation-calcination process, the CaCO<sub>3</sub>, the obtained calculations are the following.

As according to the previous equations, we have:

$$COS_{CaCO_3} = 0,00251 \text{ €/ kg CO}_2 = 2,51 \text{ €/ CO}_2 \text{ capture tonne}$$

However, these new sorbents have to be tested in order to check the reactivity and capture capacity in a higher number of cycles.

Taking into account the rest of the factors that influence in the global CO<sub>2</sub> capture cost, and making for that some assumptions, it has been estimated an approximated cost of 15,5 €

## Conclusions

The sorbent cost of the CaCO<sub>3</sub> is very cheap, carrying out at least one of the premises of how the sorbents have to be, what provokes that the big advantage of this sorbent lies in the large existent abundance of limestone. Nevertheless, the big inconvenient is that its deactivation is very quick.

This shows that the sorbents based on calcium have the potential of being economically competitive with other present capture technologies. More studies and tests have needed to improve the information about the performance of the sorbents and the equipment costs, and to give a more detailed comparison between this process and other technological alternatives.

## References

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